PATENT COOPERATION TREATY

PCT

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INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

(Chapter II of the Patent Cooperation Treaty)

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference PV/452/PCT	FOR FURTHER	ACTION	See Form PCT/IPEA/416			
International application No. PCT/CZ2005/000024	International filing dat 28.02.2005	e (day/month/year)	Priority date (day/month/year) 26.02.2004			
International Patent Classification (IPC) or no INV. C07F9/58 A61K31/663 A61P19	ational classification and 9/00	IPC				
Applicant ZENTIVA, A.S. et al						
 This report is the international preliminary examination report, established by this International Preliminary Examining Authority under Article 35 and transmitted to the applicant according to Article 36. 						
2. This REPORT consists of a total of						
3. This report is also accompanied by						
a. \square sent to the applicant and to	the International Bur	eau) a total of sheets,	as follows:			
sheets of the description, claims and/or drawings which have been amended and are the basis of this report and/or sheets containing rectifications authorized by this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions).						
Supplemental Box.	m the international ap	plication as filed, as ind	siders contain an amendment that goes licated in item 4 of Box No. I and the			
b. (sent to the International Busequence listing and/or table Relating to Sequence Listing			er of electronic carrier(s)) , containing a indicated in the Supplemental Box ructions).			
This report contains indications relations.	ating to the following i	tems:				
☐ Box No. I Basis of the repo	☐ Box No. I Basis of the report					
☐ Box No. II Priority						
☐ Box No. III Non-establishme	nt of opinion with rega	ard to novelty, inventive	step and industrial applicability			
☐ Box No. IV Lack of unity of ir	vention	, ,	otop and industrial applicability			
applicability, citat	ions and explanations	2) with regard to novelty s supporting such stater	/, inventive step or industrial nent			
☐ Box No. VI Certain documen						
	the international app					
☐ Box No. VIII Certain observation	ons on the internation	al application				
Date of submission of the demand		Date of completion of the	is report			
08.09.2005		24.05.2006				
Name and mailing address of the international preliminary examining authority:		Authorized officer	uschas Patentago			
European Patent Office - P.B. 56 NL-2280 HV Rijswijk - Pays Bas Tel. +31 70 340 - 2040 Tx: 31 68 Fax: +31 70 340 - 3016	1	Seitner, I Telephone No. +31 70 3	40-2389			

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

International application No. PCT/CZ2005/000024

_	Вох	No. I Basis of the report				
1.	. With regard to the language , this report is based on					
☐ the international application in the language in which it was filed						
	 □ a translation of the international application into , which is the language of a translation furnished for the purposes of: □ international search (under Rules 12.3(a) and 23.1(b)) □ publication of the international application (under Rule 12.4(a)) □ international preliminary examination (under Rules 55.2(a) and/or 55.3(a)) 					
2.	2. With regard to the elements* of the international application, this report is based on (replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to this report):					
	Desc	eription, Pages				
	1-12		as originally filed			
	Claim	ns, Numbers				
	1-27		as originally filed			
	Drawi	rings, Sheets				
	1-9	-	as originally filed			
	□ а	a sequence listing and/or any	related table(s) - see Supplemental Box Relating to Sequence Listing			
3.	The amendments have resulted in the cancellation of: ☐ the description, pages ☐ the claims, Nos. ☐ the drawings, sheets/figs ☐ the sequence listing (specify): ☐ any table(s) related to sequence listing (specify):					
4.	☐ This report has been established as if (some of) the amendments annexed to this report and listed below had not been made, since they have been considered to go beyond the disclosure as filed, as indicated in the Supplemental Box (Rule 70.2(c)). ☐ the description, pages ☐ the claims, Nos. ☐ the drawings, sheets/figs ☐ the sequence listing (specify): ☐ any table(s) related to sequence listing (specify):					
	* I	f item 4 applies, some	ne or all of these sheets may be marked "superseded."			

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

International application No. PCT/CZ2005/000024

Box No. V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N)

Yes: Claims

1-27

No: Claims

Inventive step (IS)

Yes: Claims

1-27

No: Claims

Industrial applicability (IA)

Yes: Claims

1-27

No: Claims

2. Citations and explanations (Rule 70.7):

see separate sheet

Re Item V

1 1

Reasoned statement with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

Reference is made to the following documents:

- D1: WO 01/56983 A (THE PROCTER & GAMBLE COMPANY) 9 August 2001 (2001-08-09)
- D2: WO 98/57967 A (DONG-A PHARMACEUTICAL CO., LTD; CHA, BONG-JIN; OH, JUN-GYO; KIM, SU-EO) 23 December 1998 (1998-12-23)
- D3: MEHTA S C: "ISSUES AND APPROACHES FOR IMPROVING THE SOLUBILITY AND BIOAVAILABILITY OF POORLY WATER SOLUBLE COMPOUNDS" BULLETIN TECHNIQUE GATTEFOSSE REPORT, GATTEFOSSE, SAINT-PRIEST,, FR, vol. 91, no. 91, 1998, pages 65-72, XP008034161 ISSN: 1149-0306

V.1. Novelty:

The present application relates to amorphous forms of Risedronic acid monosodium salt which has not been disclosed in the available prior art.

Consequently, the subject-matter of claims 1-27 is new in the sense of Article 33(2) PCT.

V.2. Inventive step:

The document D1 is regarded as being the closest prior art to the subject-matter of claim 1, and discloses Risedronic acid monosodium salt in crystalline form.

The subject-matter of claim 1 therefore differs from D1 in that the Risedronic acid monosodium salt is in amorphous form.

The problem to be solved by the present invention may therefore be regarded as the

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY (SEPARATE SHEET)

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International application No.

PCT/CZ2005/000024

provision of a further form of Risedronic acid.

The skilled person is aware of the existence of crystalline and amorphous forms and it is considered an ordinary practice to prepare different forms of active ingredients.

The Applicant provides comparative tests showing a better solubility of the amorphous form when compared to the crystalline form.

In general, it is known that amorphous forms of a drug represent the highest energy forms and are therefore used to improve the solubility/dissolution rates of insoluble drugs (see D2 and D3). Consequently, this effect cannot be regarded as unexpected.

Applicant argues that sodium risedronate, as salt of a weaker acid, converts in the environment of gastric juice, which contains HCI, to the more poorly soluble risedronic acid. In comparative tests, Applicant convincingly demonstrated that, in the case of amorphous risedronate, risedronic acid remains in the solution, while in the case of crystalline risedronate, it precipitates within a few seconds.

This effect can indeed be regarded as an unexpected effect which is associated with the novel feature over the prior art.

Therefore, the <u>subject-matter of claims 1-27 can be considered as involving an inventive step</u> (Article 33(3) PCT).

V.3. Industrial Applicability:

The present application relates to compounds which are useful for the treatment of bone diseases and the <u>subject matter of claims 1-27 is therefore considered as industrially applicable</u> (Article 33(4) PCT).